Brachydactyly type A1

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Brachydactyly type A1. ORPHA:93388

Brachydactyly type A1 (BDA1) is a congenital malformation characterized by apparent shortness (or absence) of the middle phalanges of all digits, and occasional fusion with the terminal phalanges.